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June 24, 1999



Office of Special Nutritionals (HFS-450) Center for Food Safety and Applied Nutrition Food and Drug Administration 200 C Street, SW Washington, DC 20204

RE: <u>Notification of Initial Dietary Supplement Marketing Claims</u>

Dear Sir/Madam:

Pursuant to 2 1 CFR § 101.93 this is to notify you that DCV, Inc. and its wholly owned subsidiary Legacy USA, Inc. will be marketing **a** dietary supplement containing the structure/function claims as indicated below.

1. Name and Address of Manufacturer and Distributor:

Manufacturer; Phoenix Labs i 40 Lauman Avenue Hicksville, NY 11801 Distributor:
Legacy USA, Inc.
102 Harbor City Blvd.
Melborne, FL 32901

2. <u>Text of Statement:</u>

The following statements will be made in marketing the product:

"This product:

- a) helps maintain healthy cartilage,
- b) helps the body rebuild cartilage,
- c) helps promote cartilage regeneration,
- d) encourages the production of synovial fluid, and
- e) helps maintain flexible and healthy joints."

3. <u>Description of Dietary Ingredient or Supplement:</u>

The product is a capsule containing glucosamine hydrochloride

LET 3883

4. <u>Name of Dietary Supplement:</u>

BioChoice® Flex™

5. <u>Certification of Accuracy:</u>

I, Neal Kane, Vice President of DCV, Inc. do hereby certify that the information contained in this notice is complete and accurate and that DCV, Inc. has substantiation that the statement to be made is truthful and not misleading.

Neal J. Kame